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INFORMATION DISCLOSURE STATEMENT BY APPLICANT

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Sheet

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of

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Complete if Known

Application Number	09/180,657
Filing Date	November 12, 1998
First Named Inventor	
Art Unit	1644
Examiner Name	Nolan, Patrick J.
Attorney Docket Number	021385-014500US

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FOREIGN PATENT DOCUMENTS

Examiner Initials*	Cite No. ¹	Foreign Patent Document			Publication Date MM-DD-YYYY	Name of Patentee or Applicant of Cited Document	Pages, Columns, Lines, Where Relevant Passages or Relevant Figures Appear	T ⁶
		Country Code ³	Number ⁴	Kind Code ⁵ (if known)				
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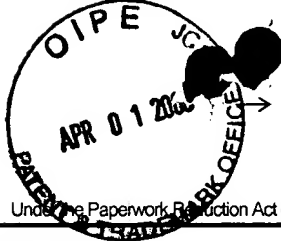
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INFORMATION DISCLOSURE STATEMENT BY APPLICANT

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Sheet 2 of 2

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Application Number	09/180,657
Filing Date	November 12, 1998
First Named Inventor	
Art Unit	1644
Examiner Name	Nolan, Patrick J.
Attorney Docket Number	021385-014500US

OTHER PRIOR ART -- NON PATENT LITERATURE DOCUMENTS

Examiner Initials *	Cite No. ¹	Include name of the author (in CAPITAL LETTERS), title of the article (when appropriate), title of the item (book, magazine, journal, serial, symposium, catalog, etc.), date, page(s), volume-issue number(s), publisher, city and/or country where published.	T ²
PN	AB	ALEXANDER, DENIS, et al; Five Related Lebanese Individuals with High Plasma Lysosomal Hydrolases: A New Defect in Mannose-6-Phosphate Receptor Recognition?; <i>American Journal of Human Genetics</i> ; 1984; pp. 1001-1014; Vol. 36, No. 5	
	AC	BROOKS, D.A., et al.; Immunoquantification of the Low Abundance Lysosomal Enzyme N-Acetylgalactosamine 4-Sulphatase; <i>Journal of Inherited Metabolic Disease</i> ; 1990; pp. 108-120; Vol. 13, No. 1	
	AD	GATTI, R., et al.; Comparative Study of 15 Lysosomal Enzymes in Chorionic Villi and Cultured Amniotic Fluid Cells. Early prenatal diagnosis in seven pregnancies at risk for lysosomal storage diseases; <i>Prenatal Diagnosis</i> ; September 1985; pp. 329-336; Vol. 5, No. 5	
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	AF	GUO, YUFENG, et al; Elevated plasma chitotriosidase activity in various lysosomal storage disorders; <i>Journal of Inherited Metabolic Disease</i> ; 1995; pp. 717-722; Vol. 18, No. 6	
	AG	KLEIMAN, FRIDA, E., et al.; Sandhoff disease in Argentina: high frequency of a splice site mutation in the HEXB gene and correlation between enzyme and DNA-based tests for heterozygote detection; <i>Human Genetics</i> ; 1994; pp. 279-282; Vol. 94	
	AH	LOVELL, KATHRYN L., et al.; Biochemical and Histochemical Analysis of Lysosomal Enzyme Activities in Caprine β -Mannosidosis; <i>Molecular and Chemical Neuropathology</i> ; 1994; pp. 61-74; Vol. 21, No. 1	
	AI	McCABE, NORAH R., et al.; Preferential Inhibition of Lysosomal Beta-Mannosidase by Sucrose; <i>Enzyme</i> ; 1990; pp. 137-145; Vol. 43, No. 3	
	AJ	O'BRIEN, JOHN S., et al.; Saposin proteins: structure, function, and role in human lysosomal storage disorders; <i>FASEB J.</i> ; 1991; pp. 301-308; Vol. 5	
	AK	PRENCE ELIZABETH M., et al.; Diagnosis of α -Mannosidosis by Measuring α -Mannosidase in Plasma; <i>Clinical Chemistry</i> ; 1992; pp. 501-503; Vol. 38, No. 4	
	AL	WHITLEY, CHESTER B., et al.; Long-Term Outcome of Hurler Syndrome Following Bone Marrow Transplantation; <i>American Journal of Medical Genetics</i> ; 15 April 1993; pp. 209-218; Vol. 46, No. 2	
	AM	YAMAGUCHI, KEIKO; Improvement of Tear Lysosomal Enzyme Levels after Treatment with Bone Marrow Transplantation in a Patient with I-Cell Disease; <i>Ophthalmic Research</i> ; 1989; pp. 226-229; Vol. 21, No. 3	

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